

**IN THE UNITED STATES  
PATENT AND TRADEMARK OFFICE**

TITLE:

Ultra Fine Dead Sea Mineral Compound and Method of Manufacture

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## **BACKGROUND OF THE INVENTION**

This application is a continuation application claiming priority to U.S. Patent Application Serial No. 09/931,453 filed August 16, 2001 and any amendments thereof.

### **1. Field of The Invention**

Applicant's invention relates to an ultra fine mineral compound and a method of processing native Dead Sea minerals into this ultra fine mineral compound that can be used to manufacture all-natural Dead Sea mineral compositions particularly compositions for use in cosmetic preparations such as bath and body products.

### **2. Background Information**

A cosmetic product is any substance or preparation intended for placing in contact with the various external parts of the human body or with the teeth or mucous membranes of the oral cavity with the intention of cleaning, perfuming, or protecting, to keep such parts in good condition, change their appearance or correct body odors. There are numerous product groups that fall within the category of cosmetic products or preparations, including but not limited to cosmetic emulsions, deodorants and antiperspirants, sunscreens, make-up preparations, hair preparations, bath products, soaps, exfoliating agents, and shaving preparations.

Cosmetic preparations are usually mixtures. A mixture is any matter consisting of two or more substances physically combined in some proportion by

1 mass. In a mixture there is no chemical reaction. Two types of mixtures are  
2 heterogenous mixtures and homogenous mixtures. A heterogenous mixture is a  
3 mixture having ingredients of different states of matter. A suspension is a  
4 heterogenous mixture in which droplets or particles are suspended in a liquid. A  
5 colloidal dispersion is a specific type of suspension in which the particles or droplets  
6 of one substance are smaller than those in suspensions , but larger than those in  
7 solutions and that have one dimension in the range of 1 to 10 nm. A homogenous  
8 mixture is a mixture having ingredients of the same states of matter. Homogenous  
9 mixtures are usually solutions which are made up of a solute dissolved in a solvent.  
10 When the solute does not remain dissolved in the solvent the mixture is in turn  
11 referred to as a heterogenous mixture.

12 Many cosmetic preparations are suspensions and more particularly colloidal  
13 dispersions. In a colloidal dispersion there is a suspension of finely divided particles  
14 in a continuous medium in which the particles do not settle out of the substance  
15 rapidly and are not readily filtered. Where the particle is a liquid droplet and the  
16 medium is a liquid, the colloid is referred to as an emulsion. If however the particle  
17 is a solid and the medium is a liquid, the colloid is referred to as a sol or gel. A sol  
18 is a colloidal dispersion of a solid in a liquid in which the particles are so small that  
19 the dispersion appears transparent while a gel is a suspension that behaves as an  
20 elastic solid or semi-solid rather than liquid.

1 Colloidal systems undergo agglomeration, or gathering into a mass, leading to  
2 a distribution of droplet size for liquid colloids. Though wetting phenomena and  
3 nonwetting colloidal factors may play a role, the agglomeration process is induced  
4 by particulate collisions arising from diffusion, as in Brownian motion, velocity or  
5 shear gradients in a liquid dispersion medium, and gravitational settling.

6 Irreversible agglomeration can be quantified using various models for  
7 repulsive or attractive electrostatic, London- van der Waals, and steric forces which  
8 affect stabilization of aqueous and nonaqueous colloidal systems. A comprehensive  
9 model of colloidal stability, the DLVO (Derjagun-Landau-Verwey-Overbeek) model  
10 has provided information regarding the roles of electrolytes, dielectric constant, and  
11 other physical quantities in colloidal systems. This theory considers the electrostatic  
12 interactions between two identically charged, suspended particles to be repulsive  
13 and to arise from the overlap of the electrical double layers associated with each  
14 particle.

15 For systems containing a soluble polymer or surfactant, molecular  
16 arrangement, thickness of the absorbed layer, temperature, and chain or segment  
17 solvation are additional critical parameters in determining the effectiveness of a  
18 dispersed agent in providing steric stabilization. If velocity or shear gradients are  
19 present, such as in mixing, and are sufficiently large, the frequency of collisions  
20 depends on the volume fraction of solids and the mean velocity gradient. Assuming  
21 that sedimentation is slow compared to the first two collision mechanisms, the

1 overall agglomeration rate is

2  $-dN/dt = k_d N^2 + k_s N$

3 where N is the particle number concentration,  $k_d$  and  $k_s$  are the respective  
4 rate constants corresponding to diffusion controlled and shear induced collision  
5 processes, and the minus sign denotes that the particle number concentration  
6 decreases with time.

7 Cosmetic emulsions, such as lotions and creams, are emulsions of water-  
8 based and oil-based phases. An emulsion is more particularly a two phase system  
9 consisting of two incompletely miscible liquids, the internal or discontinuous phase  
10 dispersed as finite globules in the other termed the continuous phase. Emulsions  
11 can be classified according which liquid is dispersed in the continuous phase. Oil in  
12 water (o/w) emulsions have oil as the dispersed phase in water as the continuous  
13 phase. In water in oil (w/o) emulsions, the water is dispersed in the oil as the  
14 continuous phase.

15 Products that produce emulsions, or emulsifiers, can be classified as ionic or  
16 nonionic according to their behavior. An ionic emulsifier is composed of an organic  
17 lipophilic group and a hydrophilic group. The hydrophilic-lipophilic balance is often  
18 used to characterize emulsifiers and related surfactant materials. The ionic types  
19 may be further divided into anionic and cationic, depending on the nature of the ion-  
20 active group. The lipophilic portion of the molecule is usually considered to be the  
21 surface active portion. Nonionic emulsifiers are completely covalent and show no

1 apparent tendency to ionize. Emulsifiers, being surface active agents, lower surface  
2 and interfacial tensions and increase the tendency of their solution to spread.

3         Mixing of cosmetic preparations is an important operation particularly in the  
4 preparation of heterogenous mixtures such as suspensions and colloids since the  
5 actual steps involved can dictate whether the particles or droplets remain suspended  
6 continuously throughout the medium for a reasonable period of time to maintain an  
7 adequate shelf life and viability of the preparation. This becomes increasingly  
8 difficult when the desire of the manufacturer is to produce cosmetic preparations  
9 that contain all natural ingredients. Natural ingredients refer to ingredients obtained  
10 from nature such as extracted directly from plants or animal products as opposed to  
11 being produced synthetically.

12         The present composition contains all natural ingredients. One of the natural  
13 ingredients incorporated into the composition of the present invention is Dead Sea  
14 minerals. Dead Sea minerals are not to be confused with sea salt or afro-salt®  
15 which has a different chemical composition. Sea salt is the compound remaining  
16 when oceanic sea water is evaporated. It contains primarily sodium and chloride  
17 and in some cases trace amounts of copper, manganese, nickel, fluorine, tin and  
18 iodine. The trace minerals can vary based upon the source of the sea water.

19 Afrosalt® is a compound of inorganic salts derived from seawater containing 45% ±  
20 31 sodium, 53% ± 3 chlorides, 3.6% magnesium, <7% sulphates, <3% calcium,  
21 <2% bromides, 0.49% ± 0.04 potassium, <0.3% iodides. The Dead Sea is a

1 unique body of water, unlike any other and has a singular chemical composition.  
2 For years it has been known that treatments administered at the Dead Sea can bring  
3 about significant remissions in diseases such as psoriasis, psoriatic arthritis,  
4 rheumatoid arthritis, and osteoarthritis. It is not known what the mode of action is  
5 of the Dead Sea minerals. It is however believed that specific ions from the  
6 minerals play a role mainly as co-factors in enzymatic regulation activities in the  
7 metabolism of healthy skin. There are indications that magnesium is a co-factor for  
8 phosphate transferring enzymes and participates in c-AMP c-GMP balancing  
9 regulation, potassium may enhance CO<sub>2</sub> transport, and calcium is thought to  
10 regulate cell membrane permeability. Zinc may play a role as a co-factor in cell  
11 proliferation enzymatic regulation.

12       Electrolytes can be absorbed into the skin from mineral rich preparations.  
13 The skin is a multilayered membrane with certain absorption characteristics which  
14 are subject to change. Corneum cell walls are involved in the semi-permeable  
15 membrane system and are responsible for the osmotic properties of the corneum.  
16 The penetration of the electrolytes through the stratum corneum occurs in between  
17 the horny cells.

18       There are models that demonstrate specific ionic absorption through the  
19 human skin barrier. Concentration is the key. When applying a cosmetic  
20 preparation, the relevant concentration is the concentration gradient between each  
21 specific dissolved ion both outside and inside the skin surface. During the

1 absorption process, a partitioning of minerals occurs from the vehicle to the skin.

2 The nature of the cosmetic preparation is significant in determining the kinetics of  
3 mineral skin penetration. Another important factor is the pH in the various  
4 microenvironments of the skin. Ions in varying valences and cations in combination  
5 with different anions penetrate to differing extents. There are major differences in  
6 the extent of skin penetration in different areas of the body.

7 The face is one of the highest absorbing areas. Exposed surface area,  
8 frequency of dermal application, skin type, skin age, temperature, and contact time  
9 should be considered. Factors involved in the percutaneous absorption of cosmetic  
10 preparations include use of other topical or systemic drugs, application parameters  
11 such as area, amount, frequency, massage; formulation such as concentration,  
12 nature of the vehicle, occlusivity, pH; formulation components such as solvents,  
13 surfactants, perfumes, dyes, inert ingredients, active ingredients, preservatives,  
14 impurities; skin damage such as abrasion, detergents, organic solvents, climatic  
15 factors; and physiological factors such as nature of the skin, anatomical site,  
16 individual factors and hydration. Assuming electrolytes can be absorbed into the  
17 skin, dermal application of mineral rich cosmetics can prove beneficial. The goal  
18 therefore has been to incorporate the beneficial properties of the Dead Sea into  
19 cosmetics.

20 Over the past few years cosmetics have been marketed that incorporate Dead  
21 Sea minerals, including body and face masks with highly viscous dispersions, lotions



1 and creams with the minerals in very low concentrations, and one phase aqueous  
2 solutions with the minerals in very low concentrations. The composition of Dead  
3 Sea minerals is very unique. The concentration of the divalent cations magnesium  
4 and calcium is very high compared with the monovalent cations, mainly sodium and  
5 potassium. In addition, the ionic strength of a solution of these minerals is very  
6 high. These two factors have a tremendous negative effect on the formation and  
7 stability of dispersions and emulsions, and strictly limit their concentration to a few  
8 percent of the weight of conventional cosmetic formulations.

9       As mentioned previously, according to the DLVO theory stabilization of  
10 dispersions of emulsions can be described as the result of the combined attraction  
11 and repulsion forces between the particles or droplets that are dispersed in  
12 continuous phases. For example, oil in water emulsions can be stabilized by the  
13 absorption of ionic surfactants onto the oil droplet surface which may become  
14 positively or negatively charged. The electric surface potential will cause repulsion  
15 between the approaching droplets. If the repulsion forces overcome the attraction  
16 forces the emulsion will be stable. The electric surface potential is strongly  
17 dependent on electrolyte concentration and on the valence of the counter ion in the  
18 solution. Therefore, the electrical repulsion is significantly reduced in systems that  
19 contain high concentrations of electrolytes in general, and divalent counterions in  
20 particular. This results in difficulties in formulating a cosmetic emulsion that  
21 contains electrolytes from the Dead Sea that is rich in magnesium and calcium

1 divalent cations at high concentrations, and will be stable for the minimum required  
2 shelf life for a cosmetic product. In addition, the high concentration of electrolytes  
3 may cause salting out and precipitation of various components of any cosmetic  
4 preparation. This may also affect the texture and the appearance of the product, its  
5 viscosity, hydrophilic-lipophilic balance, crystallization, etc.

6       The present invention provides for a chemical composition for application to  
7 the skin comprising a mixture of at least 50% processed ultra fine Dead Sea mineral  
8 particles in a continuous all natural carrier medium where in the Dead Sea mineral  
9 particles do not rapidly settle out of the carrier medium which promotes a more shelf  
10 stable product. This chemical composition takes advantage of the ionic properties of  
11 the Dead Sea minerals and contains minerals of such a fine granularity that  
12 exfoliation is not as harsh to the skin, particularly of persons suffering from severe  
13 skin disorders. In addition, the carrier medium of the present invention contains all  
14 natural ingredients and is non-comedogenic so consumers do not have to be  
15 concerned about clogged pores.

16       The problems of the prior art are overcome by the present invention by  
17 processing the Dead Sea minerals into an ultra fine mineral compound and mixing  
18 this ultra fine mineral compound with select natural ingredients using a unique swift  
19 heating, chilling and mixing technique to produce cosmetic preparations, such as  
20 body scrubs, rubs, muds, creams, lotions, and related preparations. These cosmetic  
21 preparations contain greater than 50% concentration of Dead Sea minerals but

maintain stability and preferable shelf life with a pleasant feel for the consumer.

## **SUMMARY OF THE INVENTION**

It is an object of the present invention to provide a novel method for processing native Dead Sea minerals into an ultra fine mineral compound.

Still another object of the present invention is to provide a novel ultra fine mineral compound.

It is another object of the present invention to provide a novel method for using manufactured ultra fine Dead Sea minerals to manufacture all natural Dead Sea mineral compositions.

Another object of the present invention is to provide novel Dead Sea mineral compositions.

Yet another object of the present invention is to provide novel Dead Sea mineral body scrubs, rubs, muds, creams, lotions, and related preparations.

It is still another object of the present invention to provide a novel Dead Sea mineral body scrub that remains in suspension to sustain shelf life.

It is an additional object of the present invention to provide a novel Dead Sea mineral body scrub that is made from all natural ingredients.

Still another object of the present invention is to provide a novel body scrub that has as its primary ingredient Dead Sea minerals.

It is yet another object of the present invention to provide novel Dead Sea mineral compositions that provide the optimal concentration of each specific ion in

1 skin cells.

2 Still another object of the present invention is to provide novel Dead Sea  
3 mineral compositions that provide a pleasant feel for the consumer.

4 An additional object of the present invention is to provide novel Dead Sea  
5 mineral compositions that provide the optimal delivery vehicle for the various ionic  
6 compounds.

7 Another object of the present invention is to provide novel Dead Sea mineral  
8 compositions that are not irritating to the skin.

9 In satisfaction of these and related objectives, Applicant's present invention  
10 provides an ultra fine mineral compound and a method of processing native Dead  
11 Sea minerals into this ultra fine mineral compound that can be used to manufacture  
12 all natural Dead Sea mineral compositions particularly compositions for use in bath  
13 and body products.

#### 14 **BRIEF DESCRIPTION OF THE DRAWINGS**

15 Fig. 1 is a flowchart of the organization of manufacture for the preferred  
16 embodiment of the present invention.

17 Fig. 2 is a perspective view of the conical screen mill system of the present  
18 invention manufacturing process.

19 Fig. 3 is a perspective view of the unique swift heating, chilling and mixing  
20 system of the present invention.

21 Fig. 4 is a perspective view of the tube packager of the present invention.

## **DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT**

Referring to Fig. 1, a flowchart of the organization of manufacture for the preferred embodiment of the present invention is shown. The Dead Sea mineral compositions and method for manufacture begins with the native Dead Sea minerals at step 101. The native Dead Sea minerals are typically composed of 31.0 - 35.0% magnesium chloride, 20.0-28.0% potassium chloride, 3.0-8.0% sodium chloride, 0.1-0.5% calcium chloride, 0.3-0.6% bromide, 0.05-0.2% sulfates, 0-0.3% insoluble minerals, and 32.0-40.0% water of crystallization. These values have a standard deviation of  $\pm 1\%$ . These native Dead Sea minerals are next subjected to ultra fine processing at step 102. A perspective view of the conical screen mill and impeller system 103 is illustrated in more detail in Fig. 2. In the ultra fine processing, the native Dead Sea minerals are initially held in a hopper 112, that is preferably nonmetal, dispensed to a screw conveyor 113 and transported through a primary opening 114 into a second hopper 150. From second hopper 150, the native Dead Sea minerals are dispensed into a processor 115 that contains a conical screen mill with an impeller system 103 having preferably a dual headed impeller 103b. Once in the processor 115, the native Dead Sea minerals enter into the conical screen mill 103a. Within the conical screen mill 103a, the impeller 103b forces the native Dead Sea minerals through the conical screen 103a into a third hopper 104 thereby reducing the size of the native Dead Sea minerals and forming

1 an ultra fine mineral compound. The conical screen mill with impeller system 103  
2 not only reduces the particle size of the native Dead Sea minerals, but also keeps  
3 insoluble materials, such as pieces of hard mineral or rocks, from getting through so  
4 the ultra fine minerals that are formed are essentially "rock-free". The initial  
5 granularity of the native Dead Sea minerals is typically 31.3% passing through US  
6 sieve 20 mesh, 40.2% passing through US sieve 40 mesh, 24.6% passing through  
7 US sieve 60 mesh, 2.5% passing through US sieve 80 mesh, 0.4% passing through  
8 US sieve 100 mesh, and 0.2% passing through US sieve 120 mesh, 0.4% passing  
9 through US sieve 200 mesh, and 0.3% pan. Simply, the preferred native minerals  
10 used in the present process have at a minimum 90% less than 10 mesh and 90%  
11 less than 1.7 mm size granularity. However, another grade of native minerals that  
12 can be used in the present process has a screen analysis that has at a minimum  
13 greater than 90% between 5 and 10 mesh and greater than 90% between 1.7mm  
14 and 4.0 mm size granularity. In contrast, the granularity of the ultra fine minerals  
15 consist of typically 0.0% passing through US sieve 20 mesh, 22.7% passing  
16 through US sieve 40 mesh, 29.6% passing through US sieve 60 mesh, 11.5%  
17 passing through US sieve 80 mesh, 4.9% passing through US sieve 100, 3.4%  
18 passing through US sieve 120, 9.7% passing through US sieve 200, and 18.2%  
19 pan.

1 A finer version of the minerals can also be obtained with a granularity of the  
2 minerals consisting of typically 0.0% passing through US sieve 20 mesh, 0.4%  
3 passing through US sieve 40 mesh, 16.0% passing through US sieve 60 mesh,  
4 15.0% passing through US sieve 80 mesh, 8.7% passing through US sieve 100  
5 mesh, 5.5% passing through US sieve 120 mesh, 16.2% passing through US sieve  
6 200 mesh, and 38.1% pan. The granularities for each of the minerals have a  
7 deviation of  $\pm 10\%$ . The processing step for any version of the minerals ensures  
8 that 100% will have less than 10 mesh and 100% will have less than 1.0mm size  
9 granularity.

10 At the primary opening of the processor 115, a specially fitted hose 119 is  
11 placed to regulate air into the impeller portion. This air originates from an attached  
12 air compressor 158. Before entering the impeller portion, the air passes through a  
13 pressure regulator 159 and through an air dryer 160. In addition, this air is filtered  
14 with preferably two filters, one filter 151 that removes moisture and particulates and  
15 one carbon filter 152. By maintaining a cool, dry positive pressure environment the  
16 level of heat and moisture in the process remains low enabling the chemical  
17 composition to remain natural without the necessity of adding non-natural "free  
18 flowing" or "anti-caking" agents that would alter the natural composition. A  
19 secondary opening exists between the conical screen mill and impeller system 103  
20 and third hopper 104 to allow for collection, but the remainder of this portion of  
21 processor 115 is kept closed with a cover 120, being preferably cotton or canvas,

1 to prevent the ultra fine particulate from escaping into the air and causing possible  
2 respiratory problems and to prevent unnecessary moisture from entering the ultra  
3 fine minerals and altering their chemical coordination.

4 A nuisance collection tube 105, having a collection fitting, is placed adjacent  
5 opening 153 of third hopper 104 for super fine nuisance particulate debris  
6 collection into a nuisance collection receptacle 106. The processed ultra fine  
7 minerals are next conveyed along a screw type conveyor 121 to liner bags 154  
8 within fibre drums or barrels 107 placed on a scale 155. A second nuisance  
9 collection tube 156 is placed from the screw type conveyor 121 for debris collection  
10 into a second nuisance collection receptacle 157. When 65kg of ultra fine minerals  
11 are collected, the scale 155 automatically stops the screw type conveyor 121. All  
12 of the air is removed from the liner bag 154, the liner bag 154 is lock tied, and the  
13 barrel 107 closed. The barrel 107 is then transported to a pallet (not shown) for  
14 shipment 108. Both the native Dead Sea minerals and the ultra fine processed  
15 minerals are highly hygroscopic and therefore the entire process occurs in a modified  
16 room atmosphere with a temperature no higher than 78 degrees Fahrenheit with  
17 cool, dry positive pressure.

18 Once the ultra fine minerals are prepared they can then be mixed at the  
19 mixing stage 109 into a Dead Sea mineral composition. The mixing step is  
20 illustrated in more detail with the perspective view of the swift heating, chilling and  
21 mixing system of Fig. 3. Swift heating or chilling for purposes of the present



invention is defined as heating or chilling of at least about 200 gallons in about two hours or less. Mixing performance is evaluated primarily by the physical uniformity of the ultimate composition. The elements of the mixer design are the process design, such as the fluid mechanics of the impeller, fluid regimen required by the process, scale-up, and hydraulic similarity; impeller power characteristics, including speed and diameter; and mechanical design of the mixer, such as the impeller, shafts and drive assembly.

In preparing the Dead Sea mineral composition of the present invention it is important to consider the fluid mechanics of the mixing process. Mixer power,  $P$ , produces a pumping capacity  $Q$  expressed in kg/s, and a specific velocity work term of the head  $H$  expressed in J/kg according to the formula:

$$P=QH$$

where the term  $H$  is related to the square of the velocity and therefore to fluid shear rates. If the process is dependent primarily upon the pumping capacity, the fluid velocities and the individual shear rates, both on a macro- and a micro-scale are above a certain minimum level to allow other process requirements to proceed unhindered. If the pumping capacity is increased and some of the other velocity and shear rate values are decreased below some minimum, then fluid shear stress enters into the overall design.

When mixing the Dead Sea mineral composition of the present invention which involves at its simplest level a solid and at least one liquid, the settling

1 velocity of the solid particles as well as the final viscosity of the suspension are  
2 critical factors in the process design as the ultimate goal is to obtain a composition  
3 that can be used as a cosmetic preparation with complete uniformity of the solid  
4 throughout the suspension that can be maintained not only through the mixing  
5 period, but also for a reasonable time thereafter for an adequate shelf life.

6       The process for preparing the Dead Sea mineral composition of the present  
7 invention utilizes water for many of its operations. The water originates from a local  
8 water source 168, then passes into a deionizer 169. The deionized water is then  
9 supplied to a boiler 170 and a chiller unit 123. Air is also incorporated into the  
10 system which originates from air compressor 171. The air passes from air  
11 compressor 171 through a pressure regulator 172 into an air dryer 173 to create  
12 cool dry air. The air then passes through a filter 174 and water trap 175 in process  
13 vessel 122. The process begins by heating the process vessel 122 and setting the  
14 temperature control on control panel 134 to 65 degrees Celsius to open the heat  
15 valves 124a and inject hot water from boiler 170 into the process vessel 122  
16 jacket around the surrounding inside wall of the process vessel 122. Once  
17 circulated through process vessel 122 jacket, the water is returned out valve 124d.  
18 Liquid palm oil is added to vessel 122 once the vessel 122 begins to heat. When  
19 the temperature has reached at least 35 degrees Celsius, beeswax, jojoba wax PEG  
20 120, cashew husk oil ethoxylate, and coconut oil are added to the process vessel  
21 122 to begin melting. Scrape surface 163 agitation and triple impeller 164

1 agitation are turned on from control panel 134 at slow speed to mix (these are  
2 attached to a variable speed controlled motor). When these ingredients have  
3 reached 65 degrees Celsius and are melted thoroughly together then soybean oil,  
4 olive oil, jojoba oil, and vitamin E oils (Covitol 1250 and Covi-ox) are added. The  
5 speed of the scrape surface agitation and the triple impeller agitation are increased  
6 to medium. The heating is then turned off and the flow of hot water to the process  
7 vessel jacket 122 is closed. Turbine 165 agitation is then turned on within the  
8 process vessel 122 for high speed homogenous agitation or mixing. The ultra fine  
9 minerals are added and higher speed mixing with scrape surface 163 agitation and  
10 triple impeller 164 agitation is continued. The ultra fine minerals help to reduce the  
11 temperature. A chiller unit 123 is turned on and flow valves 125 are opened to  
12 circulate chilled water. The chiller unit 123 is itself chilled by an external cooling  
13 source such as a local water supply 168. The temperature control on the process  
14 vessel 122 is then set to 45 degrees Celsius to open the cooling valves 124b to the  
15 process vessel 122. The cold water is circulated through the process vessel 122  
16 jacket and returned through valve 124c to chiller unit 123. Triple motion mixing is  
17 continued until the batch temperature reaches 45 degrees Celsius. Mixing of the  
18 batch with triple motion is continued at 45 degrees Celsius for another 15 minutes.  
19 The temperature control on control panel 134 is then set to 42 degrees Celsius and  
20 an essential oil blend is added (which is weighed and blended earlier) and mixing is  
21 continued. The batch is maintained at 42 degrees Celsius while mixing for 20-25

1 minutes. The temperature control on the process vessel 122 is then set to 40  
2 degrees Celsius and mixing is continued. When the temperature of the batch  
3 reaches 40 degrees Celsius, package tubes can then be filled. At this stage, the  
4 batch is pumped from the process vessel 122 using pump 166 with a pressure  
5 regulator 167 to a holding hopper 126 at the filling station 128 of a tube packager  
6 110. The tube packager 110 is illustrated in more detail in Fig. 4. As the level of  
7 the batch goes down in the holding hopper 126, the level sensor 176 signals the  
8 pump 166 to pump over more. The holding hopper 126 is covered except for the  
9 tube 127 that enters from the process vessel 122 due to the delicate nature of the  
10 batch product. Empty tubes are placed into a distribution station 135 and upon  
11 actuating the filling station 128 of the tube packager 110 the empty tubes are  
12 placed at station 129, oriented at station 130, filled at station 131, sealed at  
13 station 132, and trimmed into a final packaged Dead Sea mineral composition at  
14 station 133 ready for shipment 111. The tubes are preferably coextruded tubes  
15 with a barrier of protection between the mineral composition and the reactant tube  
16 surface.

17       The final Dead Sea mineral composition that can be used for cosmetic  
18 preparations such as body scrubs, rubs, muds, creams, lotions, and related  
19 preparations contains ultra fine Dead Sea minerals, palm oil, soybean oil, olive oil,  
20 jojoba oil, beeswax, essential oil blend, jojoba wax PEG 120, cashew husk oil  
21 ethoxylate, coconut oil, natural source Vitamin E oil (or d-alpha tocopherol), Vitamin

1 E oil (or natural mixed tocopherols) used as antioxidant. The preferred essential oil  
2 blend includes rosewood, lavender, chamomile, and calendula.

3 Where the final composition is a body scrub, it contains preferably  
4 approximately 51% ultra fine Dead Sea minerals, 25% palm oil, 9.0% soybean oil,  
5 5.0% olive oil, 3.0% jojoba oil, 3.0% beeswax, 1.0% essential oil blend, 1.0%  
6 jojoba wax PEG 120, 1.0% cashew husk oil ethoxylate, 1.0% coconut oil, 0.1%  
7 natural source Vitamin E oil or d alpha tocopherol, 0.05% Vitamin E oil or natural  
8 mixed tocopherols. The preferred essential oil blend for the body scrub includes  
9 0.44% rosewood, 0.34% lavender, 0.20% chamomile, and 0.02% calendula. Even  
10 with the extreme ionic character of the Dead Sea minerals, the Dead Sea mineral  
11 compositions prepared remain in suspension creating a viable cosmetic preparation  
12 that can maintain adequate shelf life and provide a more pleasant feel for the  
13 consumer.

14 Conventional methods, known to those of ordinary skill in the art of  
15 cosmetics, can be used to administer the formulation of the present invention to a  
16 user; however, the preferred administration will be by transdermal delivery.

17 Although the invention has been described with reference to specific  
18 embodiments, this description is not meant to be construed in a limited sense.  
19 Various modifications of the disclosed embodiments, as well as alternative  
20 embodiments of the inventions will become apparent to persons skilled in the art  
21 upon the reference to the description of the invention. It is, therefore, contemplated

1 that the appended claims will cover such modifications that fall within the scope of  
2 the invention.